FDA 510(k) Premarket Notification

K801768 CONTAIN X-RAY DETECTABLE LAP SPONGE

CLEARED

Aug 20, 1980
23 days to decision

K801768 · Product code: **GDY** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k801768/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Gauze/sponge, Internal, X-ray Detectable (GDY)
Date received	Jul 28, 1980
Decision date	Aug 20, 1980
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**The Buckeye Cellulose Corp.**
Location	Mchenry, IL, US
510(k) history	18 submissions · 18 cleared · 1980-1983